

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 023 884 A2

(12)

## EUROPEAN PATENT APPLICATION

(43) Date of publication:

02.08.2000 Bulletin 2000/31

(51) Int. Cl.<sup>7</sup>: A61F 13/15

(21) Application number: 99125811.2

(22) Date of filing: 23.12.1999

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU  
MC NL PT SE

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: 23.12.1998 US 220188

10.05.1999 US 309238

(71) Applicant: McNEIL-PPC, INC.

Skillman, New Jersey 08558 (US)

(72) Inventors:

- Rosenfeld, Leonard G.  
East Windsor, NJ 08520 (US)
- Foelsch, Lynn  
Hillsborough, NJ 08876 (US)

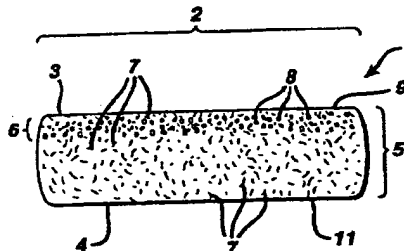
(74) Representative:

Groening, Hans Wilhelm, Dipl.-Ing.  
BOEHMERT & BOEHMERT  
Franz-Joseph-Strasse 38  
80801 München (DE)

## (54) Absorbent article with high absorbency zone

(57) The present invention is directed to an absorbent structure comprising an integral absorbent element with an upper surface and a lower surface defining a thickness therebetween. The thickness has an upper 35% containing a high absorbency zone and a lower 65%. The high absorbency zone has a first surface and a second surface separated by a zone thickness. The zone thickness is less than about 35% of the absorbent element thickness. The high absorbency zone comprises absorbent fibers and superabsorbent polymer particles. The superabsorbent polymer particles are present at a loading of no more than 75 grams per square meter and are separated from one another by the absorbent fibers. The lower 65% of the absorbent element thickness is substantially free of superabsorbent polymer particles.

FIG. 1



EP 1 023 884 A2

## Description

[0001] This is a Continuation In Part of U.S. application Serial No. 09/220,188 filed on December 23, 1998, now abandoned.

## FIELD OF THE INVENTION

[0002] The present invention relates to a novel absorbent article such as a sanitary napkin having an absorbent structure which is characterized by having an integral high absorbency zone.

## BACKGROUND OF THE INVENTION

[0003] Absorbent structures are known for inclusion in disposable absorbent articles used for absorbing body fluids and other exudates. Such absorbent structures have traditionally been made from readily available and relatively inexpensive materials such as cotton fibers, wood pulp fluff, cellulosic tissue or wadding, or other absorbent fibers. These materials have provided satisfactory absorbency of fluids both in terms of absorbency rate and overall absorbent capacity. Unfortunately, absorbent structures made from such materials may tend to collapse when wetted, thereby losing some of their void volume. Such structures may also allow absorbed fluid to be squeezed back out of the structure onto the user of the absorbent article. Furthermore, when such structures have absorbed fluid, they may present an uncomfortable wet feeling against the skin of the user.

[0004] More recently, superabsorbent polymer particles have been combined with the more traditional absorbent materials to provide structures with enhanced absorbency and retention, which may help to eliminate the problems of squeeze-out and wet surface feel. Replacement of traditional absorbent materials with superabsorbent polymer particles may also allow for absorbent products to be thinner while retaining the absorbent capacity of thicker, bulkier products. A drawback to superabsorbent polymer particles, however, is their relatively high cost compared to the more traditional absorbent materials.

[0005] Additionally, since superabsorbent polymer particles tend to swell as they absorb fluid, they may cause what is commonly known as gel-blocking. In other words, as fluid is absorbed by the particles of superabsorbent polymer, those particles swell and may form an occlusive layer of swollen superabsorbent particles. This occlusive layer then prevents the passage of additional fluid into the structure. Thus, the superabsorbent polymer particles must be properly placed within an absorbent structure to allow for this swelling and to most fully utilize their absorbent capacity. Generally, prevention of gel-blocking has been realized by mixing superabsorbent polymer particles with spacer materials, such as absorbent or nonabsorbent fibers, or by placing the superabsorbent polymer particles toward the bottom of the absorbent structure. However, although these methods of superabsorbent polymer placement may minimize gel-blocking, they do not effect the most efficient use of the superabsorbent polymer's absorbent capacity.

[0006] Therefore, what is needed is an absorbent structure with good absorbency and retention of fluid. What is also needed is an absorbent structure which will help to provide a dry feel to the skin of a user when used in an absorbent article. What is further needed is an absorbent structure with superabsorbent polymer particles spaced and placed within the structure to most fully utilize the absorbency and retention capabilities of the superabsorbent polymer particles.

## SUMMARY OF THE INVENTION

[0007] It is an object of the present invention to provide an absorbent article with good absorbency and retention of fluid which will help to provide a dry feel to the skin of a user of the article.

[0008] It is another object of the present invention to provide an absorbent structure with superabsorbent polymer particles spaced and placed within the structure to most fully utilize the absorbency and retention capabilities of the superabsorbent polymer particles.

[0009] In accordance with the present invention, there has been provided a novel absorbent structure for use in absorbent articles. The absorbent structure includes an absorbent element formed from absorbent fibers and may optionally include additional laminate layers such as one or more layers of a nonwoven fabric. The nonwoven fabric may have a lower density and a higher porosity than the absorbent element of the invention to allow for fluid acquisition and the transfer of the acquired fluid to an adjacent, higher density absorbent element. Alternatively, the nonwoven fabric may have a higher density and a lower porosity than the absorbent element to increase fluid wicking throughout the nonwoven fabric. Preferably, lower-density nonwoven fabrics are placed adjacent the body-facing surface of an absorbent element, and higher-density nonwoven fabrics are placed adjacent the garment-facing surface of an absorbent element.

[0010] The absorbent structure has peripheral edges and a center region. The center region is that portion of the

structure which is inward from the edges of the structure and which is intended accept incoming fluid when the structure is used in an absorbent articles. The peripheral edges are those portions of the structure which are adjacent its periphery.

[0011] The absorbent element has an upper surface and a lower surface defining therebetween an element thickness. The absorbent element thickness has an upper 35% and a lower 65%. The absorbent element further has an integral structure having a zone of high absorbency which comprises an integral mixture of absorbent fibers and superabsorbent polymer particles and has a first surface and a second surface separated by a zone thickness. As used herein, the terminology "integral" means a unitary structure wherein the absorbent fibers are intermeshed throughout the entire absorbent element. Thus, there are no identifiable laminate layers which are separable from other layers within the element.

[0012] The high absorbency zone is located in at least a portion of the center region of the absorbent element and within the upper 35% of the absorbent element. The first surface of the high absorbency zone may optionally be coplanar with the upper surface of the absorbent element, or alternatively, the high absorbency zone may be below the upper surface of the absorbent element, provided of course that the high absorbency zone is within the upper 35% of the thickness of the absorbent element. The thickness of the high absorbency zone may comprise up to about 35% of the thickness of the absorbent element, the remaining 65% of the absorbent element being substantially free of superabsorbent particles. Thus, the absorbent element comprises an upper high absorbency zone containing absorbent fibers and superabsorbent polymer particles located in an upper 35% of the thickness of the absorbent element and a lower absorbent region which is substantially free of super absorbent particles.

[0013] In a preferred embodiment, the superabsorbent polymer particles may be mixed with absorbent fibers within the high absorbency zone. In a most preferred embodiment, the superabsorbent polymer particles are uniformly and homogeneously mixed with the absorbent fibers within the high absorbency zone. Alternatively, the superabsorbent particles may be located within the high absorbency zone in a relatively narrow region comprising 15%, or more preferably 10%, of the absorbent element thickness. Additionally, the superabsorbent particles may be distributed within the high absorbency zone on an increasing gradient wherein the concentration of superabsorbent particles increases from the first surface of the high absorbency zone to the second surface of the high absorbency zone, or a decreasing gradient, wherein the concentration of superabsorbent particles decreases from the first surface of the high absorbency zone to the second surface of the high absorbency zone. In a most preferred embodiment the upper surface of the absorbent element is substantially free of superabsorbent particles, the high absorbency zone being slightly below the upper surface of the absorbent element wherein the upper surface including up to about 15% of the thickness of the absorbent element comprises 100% absorbent pulp fibers.

[0014] The absorbent structure is useful in absorbent articles such as sanitary napkins, diapers, incontinence articles and the like. An example of such an article comprises the absorbent element of the invention contained between a liquid-permeable body-facing layer and a liquid-impermeable barrier layer and positioned such that the body-facing layer is adjacent the upper surface of the absorbent element and the impermeable barrier layer is adjacent the lower surface of the absorbent element.

#### BRIEF DESCRIPTION OF THE DRAWINGS

[0015]

Figure 1 is a sectional view of a first preferred embodiment of the absorbent element of the invention.

Figure 2 is a sectional view of a second preferred embodiment of the absorbent element of the invention.

Figure 3 is a sectional view of a first preferred embodiment of the absorbent article of the invention.

Figure 4 is a sectional view of a second preferred embodiment of the absorbent article of the invention.

Figure 5 is a perspective view of a preferred embodiment of an absorbent article of the invention.

Figure 6 is a schematic illustration of a preferred apparatus for making the absorbent element of the invention.

Figure 7A is a detailed axial view of the particle rotary applicator of the apparatus shown in Figure 6 in the particle application phase.

Figure 7B is a detailed side view of the particle rotary applicator of the apparatus shown in Figure 6 in the particle application phase.

Figure 8A is a detailed axial view of the particle rotary applicator of the apparatus shown in Figure 6 in the recycle phase.

Figure 8B is a detailed side view of the particle rotary applicator of the apparatus shown in Figure 6 in the recycle phase.

Figure 9 is a detailed view of the gravimetric feeder and powder receivers to supply the particle rotary applicator of the apparatus shown in Figure 6.

Figure 10 is a plan view of a preferred embodiment of the absorbent element of the invention.

#### DETAILED DESCRIPTION OF THE INVENTION

[0016] Further characteristics and advantages of the invention will become clear from the following detailed description, appended drawings, and non-limiting examples.

[0017] The present invention is directed to novel absorbent articles such as, for example sanitary napkins, having an absorbent structure which includes an absorbent element having an integral high absorbency zone located adjacent an upper surface of the absorbent element. The absorbent articles will generally have a body facing, liquid-permeable cover layer, a garment-facing, liquid-impermeable barrier layer, and an absorbent structure between the body-facing layer and the barrier layer. The absorbent structure may optionally include a multi-layer laminate structure having one or more layers of nonwoven fabrics, such as fluid transfer layers in addition to the absorbent element. In a preferred embodiment, the absorbent structure comprises an upper, body facing fluid transfer layer and a lower absorbent element between the fluid transfer layer and the barrier layer. The fluid transfer layer preferably has a porosity that is greater than the porosity of the absorbent element.

[0018] The absorbent element has a center region and peripheral edges, and an upper surface and a lower surface defining therebetween an absorbent element thickness. The absorbent element thickness has an upper 35% and a lower 65%. At least a portion of the absorbent element has a high absorbency zone with a first surface and a second surface separated by a high absorbency zone thickness (hereinafter "zone thickness") comprising less than about 35% of the absorbent element thickness. The high absorbency zone is located in at least a portion of the center region in the upper 35% of the absorbent element. The first surface of the high absorbency zone may optionally be coplanar with the upper surface of the absorbent element or alternatively it may be slightly below the upper surface wherein the region between the upper surface and the first surface is substantially free of superabsorbent particles. The lower 65% of the absorbent element thickness is substantially free of superabsorbent particles.

[0019] The novel absorbent structure of the present invention is intended for use in disposable absorbent articles. These articles are adapted to be worn by the user in direct contact with the body for the purpose of absorbing body fluids and are subsequently thrown away after a single use.

[0020] Referring to Figure 1, there is shown a sectional view of a first preferred embodiment of the absorbent element 1 of the present invention. Figure 1 shows an integral absorbent element 1 with high absorbency zone 2 having a first surface 3 and a second surface 4 separated from one another by a thickness 6. The absorbent element has thickness 5 between an upper surface 23 and a lower surface 24. The high absorbency zone comprises absorbent fibers 7 and superabsorbent polymer particles 8. The superabsorbent polymer particles 8 are substantially contained within the high absorbency zone 2 in thickness region 6. It can be seen in the preferred embodiment illustrated by Figure 1 that the upper surface 23 is substantially free of superabsorbent polymer particles 8, and that the superabsorbent polymer particles 8 are separated from one another by absorbent fibers 7 in the high absorbency zone 2 throughout thickness 6.

[0021] The absorbent fibers of the present absorbent element may comprise any absorbent fiber known in the art, including without limitation, naturally occurring fibers or synthetic fibers. Examples of naturally occurring absorbent fibers are wood pulp, cotton, silk, hemp and the like, while examples of synthetic absorbent fibers include without limitation rayon fibers, individualized cross-linked cellulose fibers, acrylic fibers, and the like. A preferred absorbent fiber for the absorbent element of the invention is wood pulp fluff.

[0022] For the purposes of the present invention, the term "superabsorbent polymer" refers to materials which are capable of absorbing and retaining at least about 10 times their weight in body fluids under a 0.5 psi pressure. The superabsorbent polymer particles of the invention may be inorganic or organic crosslinked hydrophilic polymers, such as polyvinyl alcohols, polyethylene oxides, crosslinked starches, guar gum, xanthan gum, and the like. The particles may be in the form of a powder, grains, granules, or fibers. Preferred superabsorbent polymer particles for use in the present invention are crosslinked polyacrylates, such as the product offered by Sumitomo Seika Chemicals Co., Ltd. Of Osaka, Japan, under the tradename designation of SA60N Type II™, and the product offered by Chemdal International, Inc. of Palatine, Illinois, under the tradename designation of 2100A™. Although a wide range of superabsorbents are suitable for use in this invention, preferred superabsorbent particles are those well suited to being mixed at concentra-

tions of 30% or more with pulp and which do not exhibit gel blocking at these concentrations. In accordance with the present invention, sections of the high absorbency zone may contain from about 10% to about 80% superabsorbent depending on its dimensions. US Patent 5,562,646 to Goldman, which is incorporated herein in its entirety, discloses superabsorbents, which are particularly well suited to this application when the level of superabsorbent is locally greater than 60%. As disclosed more fully therein, these superabsorbents have high values of Performance Under Pressure, typically greater than 23 g/g under a confining pressure of 0.7 grams and Saline Flow Conductivity greater than  $30 \times 10^{-7}$ . These types of superabsorbents are capable of absorbing fluid, providing a dry surface feel and yet will also allow fluid to pass through the SAP region thereby minimizing any gel blocking.

**[0023]** U. S. Patent 5,601,542 to Melius, which is incorporated herein in its entirety, discloses superabsorbent particles that are particularly suitable for mixing with pulp fluff at levels of about 30%. Most preferred superabsorbents have a pressure absorbency index (as defined therein) greater than 100 and more preferably have a pressure absorbency index greater than 110 and most preferably have a pressure absorbency index greater than 120. Examples of preferred superabsorbent particles, include, but are not limited to those superabsorbent particles that are commercially available from Sumitomo Chemical Company under the tradename J550, SA60S, SA60SL, SA60SX, or from Chemdal Company under the tradename ASAP 1100, ASAP 2000, ASAP 2100, ASAP 2102, ASAP 2100A, ASAP 2300, and from Stockhausen GmbH, under the tradename Favor SXM77.

**[0024]** The absorbent elements according to the present invention may also comprise other absorbent or nonabsorbent materials, such as binders, nonabsorbent fibers, odor controlling particles, or perfumes. Examples of suitable binder materials include without limitation, ethylene vinyl acetate based latex binders, adhesives, and thermally fusible fibers, such as bicomponent fibers. Examples of suitable nonabsorbent fibers include without limitation, polyester fibers, polyolefin fibers, and bicomponent fibers.

**[0025]** Absorbent elements according to the present invention are commonly formed by air-laying the fibers and superabsorbent polymer particles. A preferred method of forming the absorbent element of the invention involves first forming pulp fluff from a pulp board in a hammer mill or similar equipment designed to fiberize or separate and "open" the pulp fibers in the board. The separated pulp fibers are then entrained in an air stream and deposited on a foraminous surface to form a pulp batt or pad. The pulp batt or pad thus formed is a collection of individual fibers in a very loose configuration. The fibrous batt is substantially uncompressed, leaving spaces between the fibers that comprise the batt. Superabsorbent polymer particles that are added to the loose batt fall into these spaces between the fibers. The superabsorbent polymer particles may be added to a portion of the air-entrained fibers for deposition substantially throughout the thickness of the high absorbency zone. Alternatively, the superabsorbent polymer particles may be deposited directly onto a formed pulp batt at a desired point in the pulp deposition process to ensure that the superabsorbent particles are located at the desired narrow zone within the thickness of the structure. In the former case, the particles are mixed with pulp fibers throughout the high absorbency zone of the integral absorbent structure. In the latter case, the particles fall into the spaces between the fibers to form a fairly concentrated high absorbency zone within the integral absorbent structure with fibers separating the particles. In either case, the particles are separated by fibers. Finally, in a preferred embodiment, pulp fibers are laid over the top of the high absorbency zone so that the upper surface of the absorbent element is substantially free of superabsorbent polymer particles. In all cases, however, the superabsorbent polymer particles are substantially separated from one another by intermeshed pulp fibers within the high absorbency zone to maintain the integral configuration of the absorbent element.

**[0026]** The high absorbency zone located adjacent the upper surface of the absorbent element may extend across the entire upper surface of absorbent element or alternatively, may be confined to a particular localized region of the absorbent element, such as for example, being located solely in a central region and spaced inwardly away from the longitudinal edges or transverse end regions of the absorbent element. Alternatively, in another embodiment, the high absorbency zone may comprise multiple discrete regions substantially separated from one another. In accordance with this embodiment, a first high absorbency zone is located in a center region of the absorbent element and one or more longitudinally or transversely oriented high absorbency zones may be located adjacent to the longitudinally extending sides or across the transverse end regions in a spaced apart relationship to one another.

**[0027]** The thickness of the absorbent structure may be uniform throughout the expanse of the absorbent element or, for the purpose of specific fit, flexibility and absorbency requirements, the absorbent structure may have a tapered profile wherein certain areas of the structure, such as a central region, are thicker than other areas.

**[0028]** As shown in Figures 2 and 4, the high absorbency zone may be surrounded by boundaries, such as densified channels. In this embodiment, the high absorbency zone comprises about 35% of the thickness of the absorbent element and is completely contained in a centrally located region between the channels. While the high absorbency zone may extend into the densified channels, it is preferred that the channels be substantially free of superabsorbent particles. The boundaries may also comprise other structural elements, such as raised areas measuring a greater thickness or caliper than the surrounding areas; repellent-treated areas; embossed or depressed areas measuring a lesser caliper or thickness than the surrounding areas; colored areas having inks or other coloring agents printed thereon or being otherwise treated to exhibit a color that is visually perceptible as different from the color of the surrounding areas;

or the edges of the absorbent structure. Alternatively, the top surface of the structure may be partially covered by a film or other impermeable material, leaving only a central opening uncovered. In such a case, this central opening would be the high absorbency zone.

[0029] An important feature of the present invention is that the entire absorbent element, including the high absorbency zone, be an integral structure, wherein the absorbent fibers that comprise the absorbent element are continuously intermingled with the absorbent fibers that comprise the high absorbency zone, and no discernible laminate or separate structure is present. In other words, the superabsorbent-containing high absorbency zone is not present as a separate layer within the absorbent element; rather, it is simply a region within the absorbent element. An advantage of this integral type of structure is that it remains a single whole structure during the absorption of fluid, and it is not subject to delamination or gapping as fluid is absorbed. Such gapping or delamination disrupts the fluid transport and distribution capabilities of the structure and may diminish its overall absorbency. An integral absorbent structure also allows for a simpler process of absorbent article construction.

[0030] The absorbent element shown in Figure 2 is an alternative embodiment of the present invention. Figure 2 shows an integral absorbent element comprising a high absorbency zone 2 with a first surface 3 and a second surface 4 separated from one another by a thickness 6. The absorbent element has thickness 5 and high absorbency zone 2 with zone thickness 6 adjacent the first surface. The absorbent element comprises absorbent fibers 7 and superabsorbent polymer particles 8. The superabsorbent polymer particles 8 are substantially contained within the high absorbency zone thickness 6. It can be seen in the embodiment illustrated by Figure 2 that the upper surface 23 is substantially free of superabsorbent polymer particles 8, and that the superabsorbent polymer particles 8 are mixed with absorbent fibers 7 in the high absorbency zone thickness 6. Figure 2 also shows the presence of two densified channels 12, which may be compressed into the absorbent element after its formation. Each channel 12 has inner edges 13 and a lowermost portion or bottom 14. The channels define the high absorbency zone therebetween. Side portions 15 of the element include those portions of the element that are outboard of channels 12, or those portions of the article that are between the top 16 of the channel and the outside edge 17 of the element. These side portions 15 may contain superabsorbent or they may be free of superabsorbent. Each side portion 15 shown in Figure 2 comprises a top surface 18 and a bottom surface 19 separated from one another by a side portion thickness 21. The side portion thickness 21 may be substantially equivalent to the high absorbency zone thickness 5, as shown in Figure 2; however, side portion thickness 21 may be substantially more or less than high absorbency zone thickness 6.

[0031] The side portion thickness 21 may comprise a second high absorbency zone thickness 22 adjacent the top surface 18. The second high absorbency zone 22 comprises 35% of the side portion thickness 21. As shown in Figure 2, each side portion 15 comprises absorbent fibers 7 and superabsorbent polymer particles 8, wherein the superabsorbent polymer particles 8 are substantially contained within the second region 22.

[0032] The absorbent article 30 shown in Figure 3 is a sectional view of a first preferred embodiment of the absorbent article of the present invention. The absorbent article 30 of Figure 3 has an integral absorbent element comprising a high absorbency zone 2 with a first surface 3 and a second surface 4 separated from one another by a thickness 6. As discussed above, the high absorbency zone is integrally formed with the absorbent element and thus the first and second surfaces 3 and 4 are not, per se, identifiable surfaces. Rather, they are marked by an absence of any superabsorbent polymer particles. The absorbent element thickness 5 has a high absorbency zone thickness 6 in the upper 35%. The absorbent element comprises absorbent fibers 7 and superabsorbent polymer particles 8. The superabsorbent polymer particles 8 are substantially contained within the high absorbency zone thickness 6. It can be seen in the preferred embodiment illustrated by Figure 3 that the upper surface 23 is substantially free of superabsorbent polymer particles 8, and that the superabsorbent polymer particles 8 are mixed with absorbent fibers 7 in the high absorbency zone thickness 6. The integral absorbent element is overlaid with a fluid transfer layer 32 and positioned between a body-facing layer 33 and a barrier layer 34 such that the upper surface 23 is adjacent the fluid transfer layer 32, which is adjacent the body-facing layer 33. The body-facing layer 33 and the barrier layer 34 are joined to one another around the periphery of the absorbent element to form what is commonly known as a flange seal.

[0033] The absorbent article 40 shown in Figure 4 is a sectional view of a second preferred embodiment of the absorbent article of the present invention. The absorbent article 40 of Figure 4 has an integral absorbent element comprising a high absorbency zone 2 with a first surface 3 and a second surface 4 separated from one another by a thickness 6, essentially as shown in Figure 2 and described in detail above. As shown in Figure 4, the integral absorbent element of Figure 2 is positioned between a body-facing layer 33 and a barrier layer 34 such that the upper surface 23 is adjacent the body-facing layer 33. The body-facing layer 33 follows the shape of the upper surface to line the inner edges 13 of the channels. The body-facing layer 33 may be connected or secured to the bottom 14, or lowermost portion, of the channel 12, as shown in Figure 4.

[0034] The body-facing layer 33 and the barrier layer 34 are joined to one another around the periphery of the absorbent element to form what is commonly known as a flange seal 36.

[0035] Although the articles 30 and 40 shown in Figures 3 and 4 respectively have the body-facing layer 33 and the barrier layer 34 joined together by a flange seal 36, this is for illustrative purposes only. The presence of a flange seal

is not necessary to achieve the benefits and advantages of the invention. Alternatively, the body-facing layer of the absorbent article may be wrapped completely around the absorbent element and overlapped and sealed to itself on the underside of the article. The barrier layer may be present either between the absorbent element and the overlapped portion of the body-facing layer, or on the outer surface of the overlapped portion of the body-facing layer. Other methods of securing a body-facing layer and a barrier layer to an absorbent article structure will be apparent to those who are familiar with the construction of absorbent articles.

**[0036]** The body-facing cover layer may comprise any soft, flexible, porous material which allows fluid to pass there-through, and may, for example, be comprised of a nonwoven fibrous sheet or an apertured or perforated plastic film. Examples of suitable nonwoven fibrous sheets include without limitation, carded webs, spunbond webs, meltblown webs, random-laid webs, and the like. The fibers comprising such webs may comprise polyester, polyethylene, polypropylene, rayon, or combinations of these. The webs may further comprise bonding agents, surfactants, or other treatments. A preferred material for the body-facing layer of the invention is a homogeneous blend of high denier polypropylene staple fibers and low denier polypropylene staple fibers. The high denier staple fibers and the low denier staple fibers preferably differ by 2 denier, where the low denier staple fibers preferably have a denier of about 3 and the high denier staple fibers preferably have a denier of about 5. The high denier staple fibers are present in the non-woven fabric in an amount of from 40 to 60 weight percent. The low denier staple fibers are present in the non-woven fabric in an amount of from 40 to 60 weight percent based on the total weight of the non-woven fabric.

**[0037]** The barrier layer is a liquid-impermeable layer, and may comprise any flexible material that prevents the transfer of fluid but does not necessarily prevent the passages of gases. Commonly used materials are polyethylene or polypropylene films. Other suitable polymeric film materials that may be used as impermeable barriers include, but are not limited to polyesters, polyamides, polyethylene vinyl acetate, polyvinyl chloride, and polyvinylidene chloride, and the like and combinations thereof. Co-extruded and laminated combinations of the foregoing, wherein such combinations are permitted by the chemical and physical properties of the film, may be used. Fluid impermeable foams and repellent treated papers may also be used. Films that are fluid barriers, but permit gases to transpire, i.e., "breathable films", may be used. These may be chosen from polyurethane films and from micro-porous films, where micro-porosity is created by ionizing radiation or by leaching out of soluble inclusions using aqueous or non-aqueous solvents. Fabrics whose surfaces have been made repellent or whose pores are small by virtue of close packing of fibers, or whose pores have been reduced in size by closing off large liquid admitting pores, may also be used alone, or together with breathable films, as breathable barriers.

**[0038]** A suitable backing sheet material can be an opaque polyolefin, e.g., polyethylene, web impermeable to body fluids and about 0.001 inch thick. Another suitable sheet material for this purpose is a polyester, e.g., polyethylene terephthalate, web having a thickness of about 0.0005 inch.

**[0039]** A preferred apparatus for making the absorbent structure of the invention is illustrated in Figure 6. With reference to Figure 6, the absorbent element of the present invention may be prepared according to the following method. While any of the absorbent fibers as previously discussed may be used to form the absorbent element, for purposes of illustration, wood pulp fibers are used to describe the preferred apparatus. The wood pulp of the absorbent element is supplied in raw material form as a compressed sheet, or pulp board 50, that is wound on a roll. The pulp unwind 51 allows the board to be fed into a pulp mill 52, where a high speed hammer rotor opens the board into substantially individual wood pulp fibers of about 2.5 mm average length, commonly known as pulp fluff or ground wood pulp. Air is pulled through the pulp mill and the adjacent forming chamber 53 by a forming wheel vacuum 54. This air conveys the pulp fluff to a forming wheel 55 and into a mold 56. The molds 56 are cavities in the forming wheel surface spaced around the circumference of the forming wheel 55. The bottom of the molds comprises a porous screen to allow the air to be pulled through the molds, leaving the pulp fluff deposited on the screen.

**[0040]** The molds are mounted on the forming wheel which rotates clockwise. When the molds first enter the forming chamber at position A, they are empty. In the Initial Fiber Deposition Zone 57, 100% pulp fibers are deposited on the bottom of the molds 56. The thickness of pulp deposited in the Initial Fiber Deposition Zone comprises 5% to 25% of the final thickness of the absorbent element, and it acts as a filter to hold the granular superabsorbent polymer powder that will be deposited in the mold. The boundaries of the Initial Fiber Deposition Zone 57 are formed by the left side of the forming chamber 53 and the left side of the seal for the Powder Application Zone 58.

**[0041]** The Particle Application Zone 58 comprises a rotary particle applicator valve 59 that dispenses a predetermined amount of particles into the pulp fluff in each mold. The particles are applied in a pattern that is phased with the molds to form the high absorbency zone, and in this embodiment, such that the high absorbency zone is substantially centered in the mold. Preferably, the high absorbency zone is spaced at least 3 mm inwardly from the peripheral edges of the mold. Most preferably, the high absorbency zone is spaced at least 7 mm inwardly from the peripheral edges of the mold and therefore, also at least 7 mm from the peripheral edges of the absorbent element contained therein. Although the high absorbency zone shown in this embodiment is generally rectangular, the shape of the high absorbency zone is not limited to rectangular. Any shape of high absorbency zone may be used and one of ordinary skill in the art will discern that varying high absorbency zone shapes may be desirable or even preferable for varying shapes and

types of absorbent elements.

[0042] The molds then enter the Final Fiber Deposition Zone 60 wherein additional pulp covers the high absorbency zone thereby forming an integral absorbent element. The molds are slightly overfilled with pulp, and two scarfing brushes 61 are used to make the pulp even with the top of the mold. The absorbent elements are then vacuum transferred out of the molds onto the Vacuum Transfer Drum 62, from which they may then be transferred to another forming station for incorporation into absorbent products.

[0043] Figures 7A, 7B, 8A, 8B, and 9, illustrate the operation of the Rotary Particle Application Valve 59 in the Particle Application Zone 58 in greater detail.

[0044] Figures 7A and 7B show an axial view and a side view respectively of the Rotary Particle Application Valve 59 in the particle application phase. In order to achieve a precise pattern of particles on each mold 56, Rotary Particle Application Valve 59 is used as a means to start and stop the flow of particles to the molds 56. Particles are delivered to the Rotary Particle Application Valve 59 in the Particle Application Zone 58. Preferably, the particles are delivered via a gravimetric feeder, such as a loss-in-weight (LIW) screw feeder 70 to accurately control the particle weight applied to each mold 56. The particle supply source 72 is located outside of the Particle Application Zone. The discharge end of the screw feeder 70 is located within a stationary funnel 74 of the Rotary Particle Application Valve. The stationary funnel is housed within the rotor 76 of the Rotary Particle Application Valve. The rotor 76 comprises at least one rotor slot opening 78. The widths of the stationary funnel 74, the rotor slot opening 78, and the discharge end of the screw feeder 70 are matched to the width of the desired pattern of particles to be formed on each mold, which determines the shape of the high absorbency zone. As the rotor 76 turns, and the rotor slot opening 78 aligns with the discharge end of the screw feeder 70, the superabsorbent polymer particles 8 contained in the stationary funnel drop by gravity onto the mold 56. The forming wheel vacuum 54 assists with the drawing of the particles downward onto the mold 56. Preferably, a portion of the screen bottom of each mold 56 is also masked such that an opening remains having the desired pattern shape. This selective masking of the pulp molds enhances the accurate and precise placement of the particles within the pulp mold. The length of the rotor slot 78 dictates the length of the pattern of particle forming the high absorbency zone 2 of the absorbent element 1.

[0045] Figures 8A and 8B show an axial view and a side view respectively of the Rotary Particle Application Valve 59 in the recycle phase. In Figures 8A and 8B, the forming wheel 55 is shown in a position such that the Rotary Particle Application Valve 59 is located over a portion of the wheel between two molds 56. It is desired to prevent the deposition of particles over this portion of the wheel, since any particles dispensed in this position are essentially wasted and serve only to contaminate the area surrounding the apparatus. The recycle phase shown in Figures 8A and 8B prevents the problems associated with unwanted disposition of particles by recycling the particles. When the Rotary Particle Application Valve 59 is in the recycle phase as shown, the position of the rotor 76 under the stationary funnel 74 prevents the passage of particles; i.e., the rotor slot 78 is in a closed position. Particles exiting the screw feeder 70 hit the inside diameter of the rotor. A rotor vacuum port 80 in the side of the rotor opens to the rotor inside diameter through a series of recycle holes 82 in the rotor and pulls the particles out of the rotor and into a recycling tube 84. As shown in Figure 9, the particles are then conveyed by air through the recycling tube 84 to a recycle receiver 102 that will eventually provide the particles back to the screw feeder 70 for reuse. Also shown in Figure 9 is a detailed view of the particle supply source 72, including both the virgin supply reservoir 104 and the recycle receiver 102.

[0046] Figure 10 shows a simplified plan view of an absorbent element 1 made by the process, including the high absorbency zone 2. The length and width of the high absorbency zone are determined by the length and width of the rotor slot opening 78.

#### EXAMPLE

[0047] An absorbent element was formed by a vacuum forming process as illustrated in Figure 6 and as described herein, in which a pulp board was defiberized and dispersed into individual fibers in a hammer mill. The pulp fibers were entrained in a stream of air and enough of the fibers were deposited into a shaped screen-bottomed mold to cover the screen with about a one (1) millimeter thickness of pulp fluff. A predetermined amount of Chemdal 2000™ superabsorbent polymer particles was then deposited in a 35 millimeter wide stripe along a central portion of the mold. The amount of superabsorbent polymer particles (SAP) was either 0 grams, 0.2 gram, 0.4 gram, or 0.7 gram per mold. Since the deposition of superabsorbent polymer particles per mold was substantially evenly disposed along a 35 mm wide stripe on the absorbent structure, and since the superabsorbent polymer particles tended in general to be somewhat more dense than pulp fluff fibers, the superabsorbent polymer particles tended to lodge in the spaces between pulp fluff fibers as they were dispensed onto the 1 mm thickness of pulp fluff. Finally, additional pulp fibers were added to the mold to form an absorbent element with a total pulp weight of 8.0 grams and a total weight of superabsorbent polymer particles as provided in Table 1.

[0048] The element was removed from the mold and oriented so that the surface of the element which had contacted the screen was the upper or top surface of the element. The resulting element had a generally oblong shape with



a length of about 200 millimeters and a width of about 60 millimeters. Each absorbent element was between 7 mm and 20 mm in thickness with a density of less than 0.07 g/cc.

[0049] Each absorbent element was then placed between a cover layer comprising a homogeneous blend of 50% of 5 denier polypropylene staple fibers and 50% of 3 denier polypropylene staple fibers, and a barrier layer of 0.001 inch thick polyethylene film with the cover layer being placed on the first surface of the element, to form an absorbent article. The cover and barrier were sealed to one another around the edges of the absorbent element to form a flange seal. Densified side channels were then formed in the upper (cover layer) surface of the article by heating and compressing the cover/absorbent element/barrier assembly in selected areas to form densified channels similar to those shown in Figure 5. The density of the channels was at least 0.5 g/cc.

[0050] The articles were then tested for strikethrough times and rewet amounts. Strikethrough times indicate the amount of time that is required for each of six 2 cc aliquots of test fluid to penetrate the cover, where lower numbers indicate faster fluid penetration and thus, less gel-blocking. Rewet amounts indicate the amount of absorbed test fluid that may be squeezed out through the cover of the product when pressure is applied. Lower rewet amounts indicate that the absorbed fluid is being retained well, thus leading to a drier feeling product in use.

[0051] The strikethrough time was measured by the following method. A 2.54 cm thick plate with a 2cm round hole was placed over the cover of the article with the hole aligned with the center of the article. A 2 cc dose of synthetic menstrual fluid was deposited in the hole and the time required for it to absorb into the article (i.e., penetration time) was measured. Five additional 2 cc doses were added at 5 minute intervals and the penetration time was measured for each one. The penetration times for the first and sixth doses are shown in Table 1. The increases in penetration time for the sixth dose with 0.4 and 0.7 gram SAP addition are evidence of gel blocking.

[0052] For the rewet test, a 2.54 cm thick orifice plate with a centrally located 3.8 cm by 1.9 cm elliptical aperture was placed over the article with the aperture aligned with the center of the article. A 7 cc dose of synthetic menstrual fluid was deposited in the aperture and allowed to penetrate the article. After 15 minutes a pre-weighed stack of four 2-inch X 4-inch sheets of Whatman #1 filter paper was placed over the article sample. A 2-inch X 4-inch 2.2 kg weight was placed over the filter paper for three (3) minutes. The filter paper was then removed and weighed. The amount of fluid picked up by the filter paper was calculated and reported as the rewet amount.

[0053] The test fluid used for the penetration test and the rewet test may be any synthetic menstrual fluid having a viscosity of approximately 30 centipoise. The results of the tests are indicated in Table I below.

TABLE I

35 mm Wide strip of SAP along length of absorbent element 8.0 Gram Pulp				
Amt. SAP (grams)	Amt. SAP (gsm)	Rewet Value (grams)	Penetration (1 <sup>st</sup> dose - sec)	Penetration (6 <sup>th</sup> dose - sec)
0	0	0.027	6	13
0.2	30	0.019	6	13.0
0.4	60	0.021	6	24
0.7	103	0.025	6.5	31

[0054] As can be seen from the data in Table I, the formation of an integral high absorbency zone comprising super-absorbent polymer improved the rewet amounts, with the 30 gsm loading of superabsorbent leading to the best (lowest) rewet amount. Slightly higher rewet was measured at 60 gsm. At 103 gsm, the rewet was not improved substantially compared to the element without SAP, and the penetration time was sharply increased. Therefore, it was discovered that for the absorbent element of the present invention, a superabsorbent loading of between 30 -55 gsm is preferred for improving rewet amount without gel-blocking. Some rewet benefit may also be obtained at SAP add-ons of up to 75 gsm.

[0055] The specification and embodiments above are presented to aid in the complete and non-limiting understanding of the invention disclosed herein. Since many variations and embodiments of the invention can be made without departing from its spirit and scope, the invention resides in the claims hereinafter appended.

## Claims

1. An absorbent structure having peripheral edges and a center region and comprising an integral absorbent element (1) having an upper surface and a lower surface defining therebetween an absorbent element thickness (6), the

absorbent element thickness (6) having an upper 35% and a lower 65%, the absorbent element (1) further comprising a high absorbency zone (2), the high absorbency zone (2) having a first surface (3) and a second surface (4) separated by a zone thickness (6) comprising less than about 35% of the absorbent element thickness (6) and being located in at least a portion of the center region in the upper 35% of the absorbent element (1), the high absorbency zone (2) comprising absorbent fibers (7) and super-absorbent polymer particles (8), the superabsorbent polymer particles (8) being present at a loading of no more than 75 grams per square meter, the superabsorbent polymer particles (8) being substantially separated from one another by absorbent fibers (7), and wherein the lower 65% is substantially free of superabsorbent polymer particles (8).

2. The absorbent structure of claim 1 wherein the superabsorbent polymer particles (8) are mixed with absorbent fibers (7) within the high absorbency zone thickness (6).

3. The absorbent structure of claim 2 wherein the superabsorbent polymer particles (8) are uniformly mixed with absorbent fibers (7) within the high absorbency zone thickness (6).

4. The absorbent structure of claim 1 wherein the superabsorbent polymer particles (8) are present at a loading of between 30 and 55 grams per square meter.

5. The absorbent structure of claim 1 wherein the absorbent element upper surface (23) is substantially free of superabsorbent polymer particles (8).

6. The absorbent structure of claim 1 having at least two spaced apart densified channels (12) and wherein the high absorbency zone is confined between the at least two densified channels (12).

7. The absorbent structure of claim 1 wherein the high absorbency zone comprises a substantially rectangular pattern of superabsorbent polymer particles (8) at least partially contained in the center region.

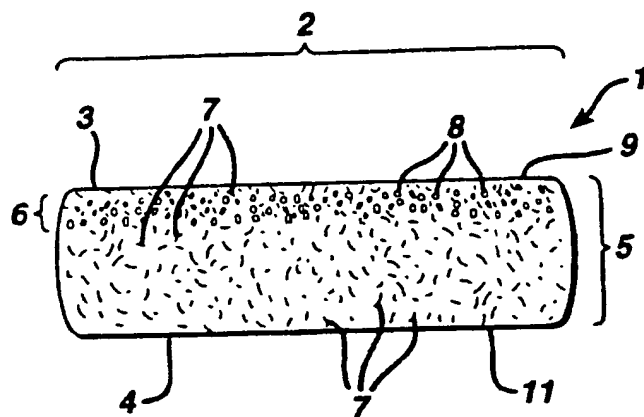
8. The absorbent structure of claim 1 wherein the high absorbency zone thickness (6) comprises about 15% of the absorbent element thickness (5), the high absorbency zone being located within an upper 25% of the absorbent element thickness.

9. The absorbent structure of claim 1 wherein the high absorbency zone thickness comprises about 10% of the absorbent element thickness, the high absorbency zone (2) being located within an upper 20% of the absorbent element thickness (5).

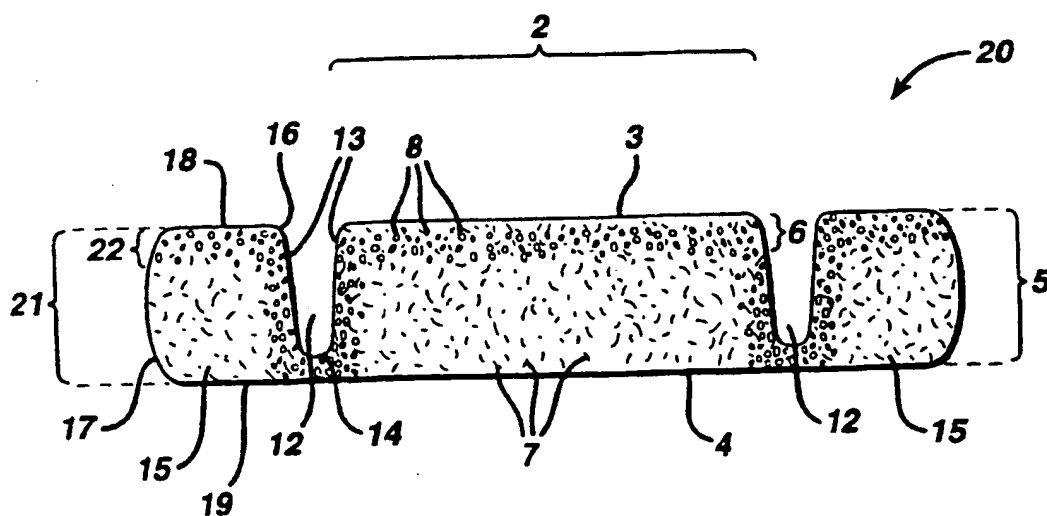
10. An absorbent article (40) adapted to be worn in a crotch portion of a user's undergarment, the article comprising:

- a liquid permeable body-facing layer (33);
- a liquid impermeable barrier layer (34); and
- an absorbent structure (2) positioned between the bodyfacing layer (33) and the barrier layer (34) such that the first surface (3) is adjacent the body-facing layer (33), the absorbent structure having peripheral edges and a center region and comprising an integral absorbent element (2) having an upper surface (3) and a lower surface (4) defining therebetween an absorbent element thickness (6), the absorbent element thickness (6) having an upper 35% and a lower 65%, the absorbent element (2) further comprising a high absorbency zone (22), the high absorbency zone (22) having a first surface and a second surface separated by a zone thickness (6) comprising less than about 35% of the absorbent element thickness (6) and being located in at least a portion of the center region in the upper 35% of the absorbent element (2), the high absorbency zone (22) comprising absorbent fibers (7) and superabsorbent polymer particles (8), the superabsorbent polymer particles (8) being present at a loading of no more than 75 grams per square meter, the superabsorbent polymer particles (8) being substantially separated from one another by absorbent fibers (7), and wherein the lower 65% is substantially free of superabsorbent polymer particles (8).

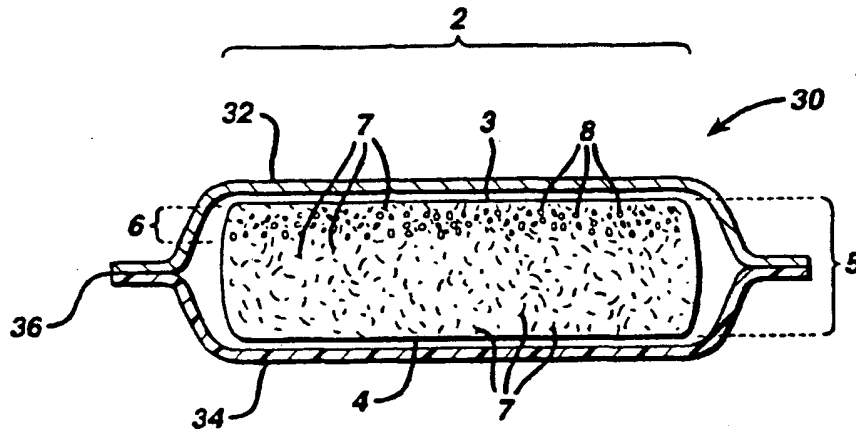
**FIG. 1**



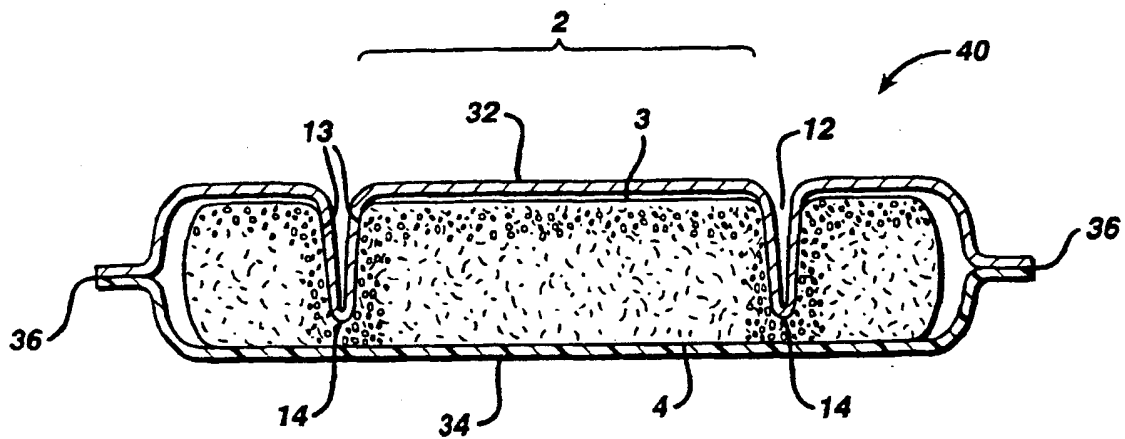
**FIG. 2**



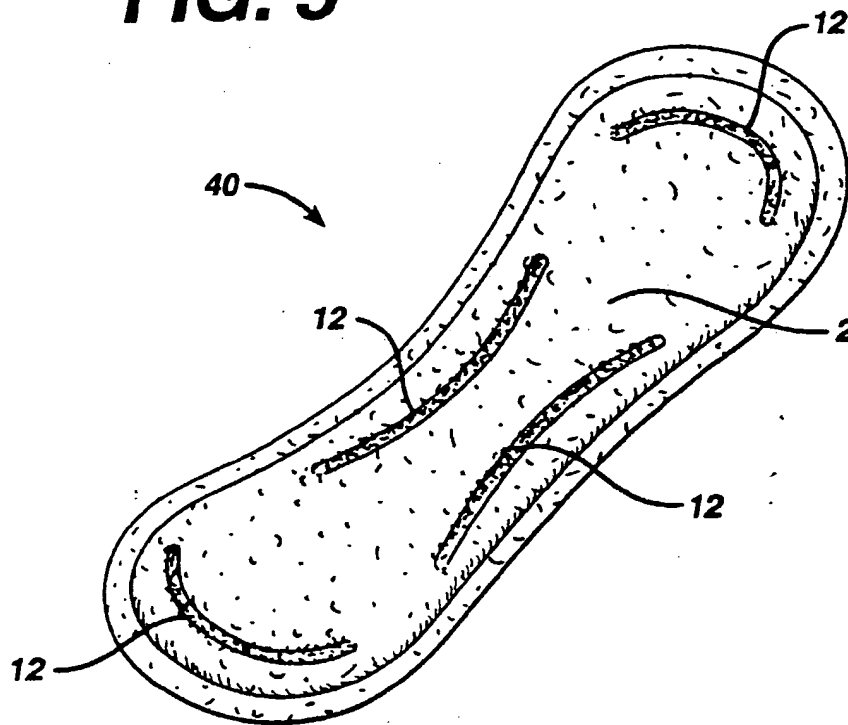
**FIG. 3**



**FIG. 4**



**FIG. 5**



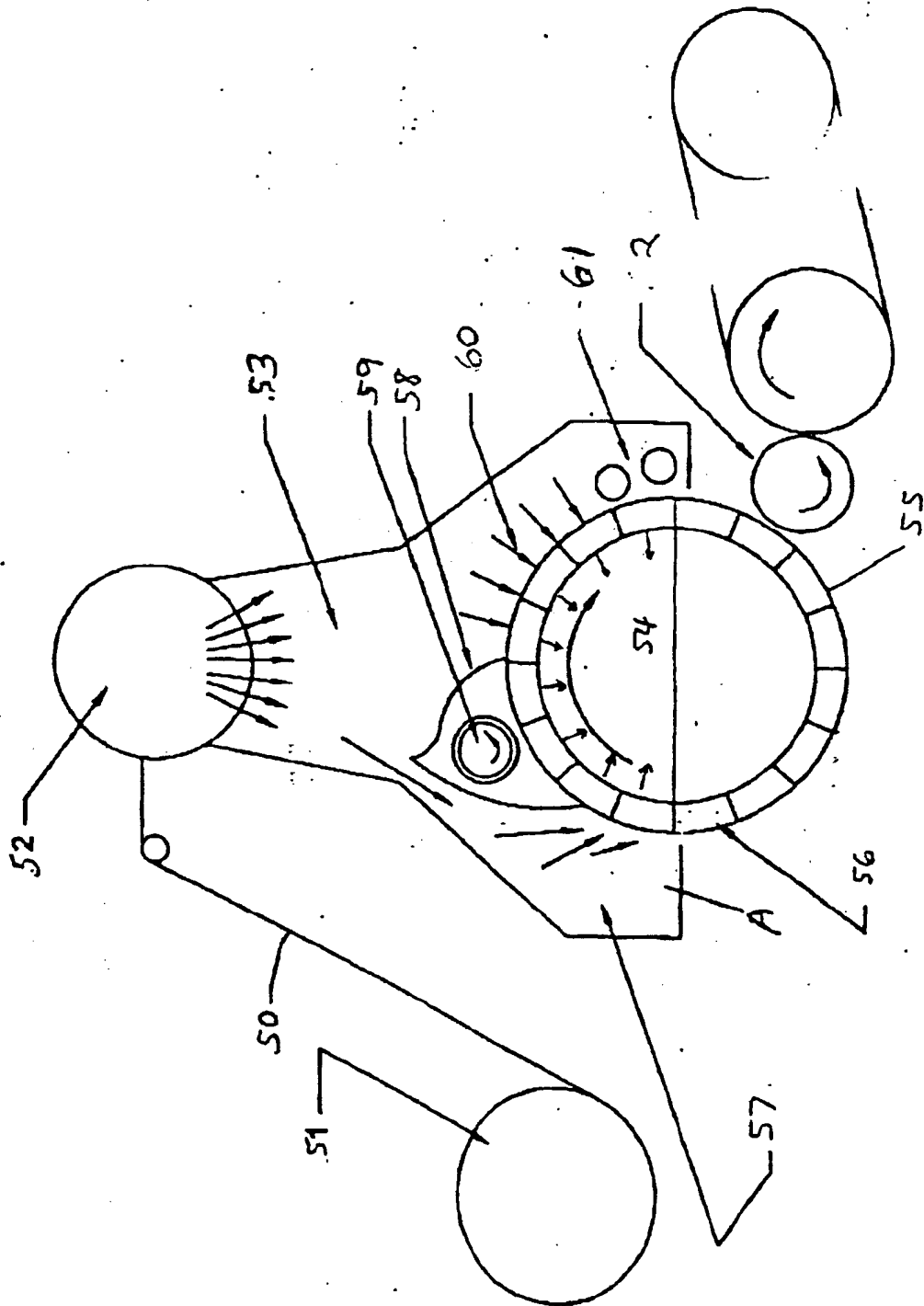


Figure 6

POWDER APPLICATION PHASE

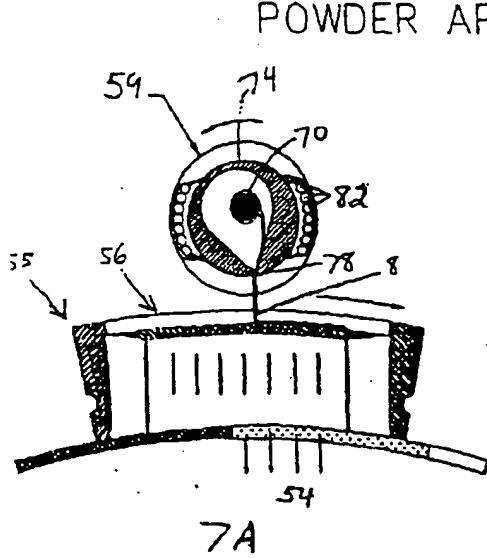
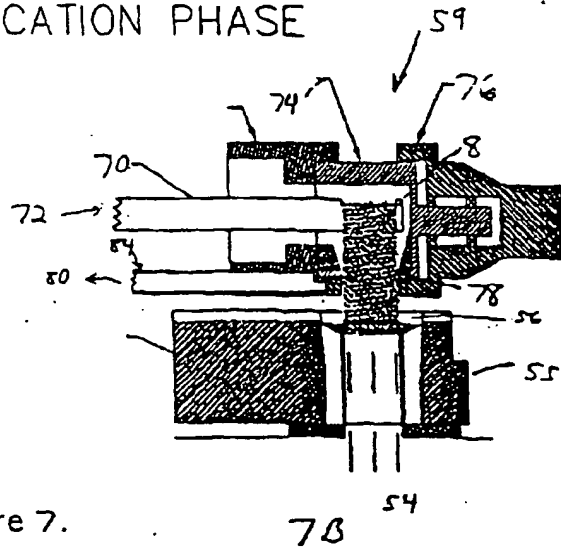


figure 7.



7B

RECYCLE PHASE

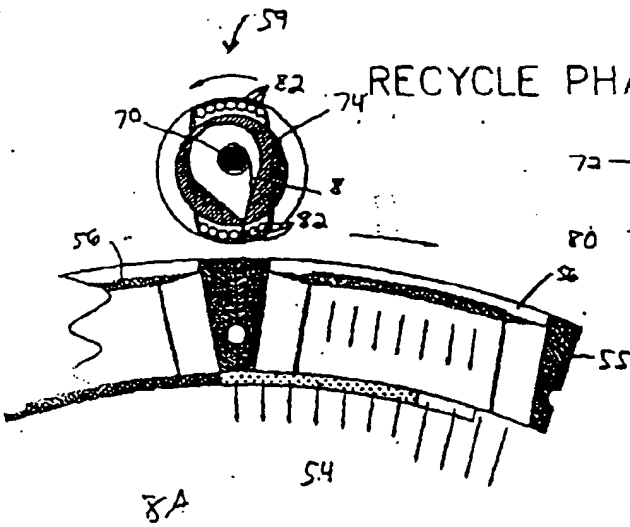
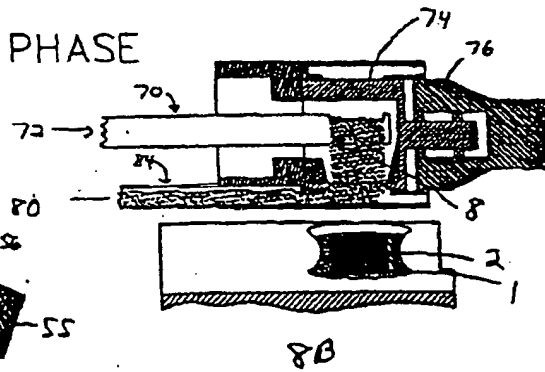


figure 8.



8B

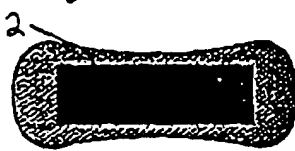
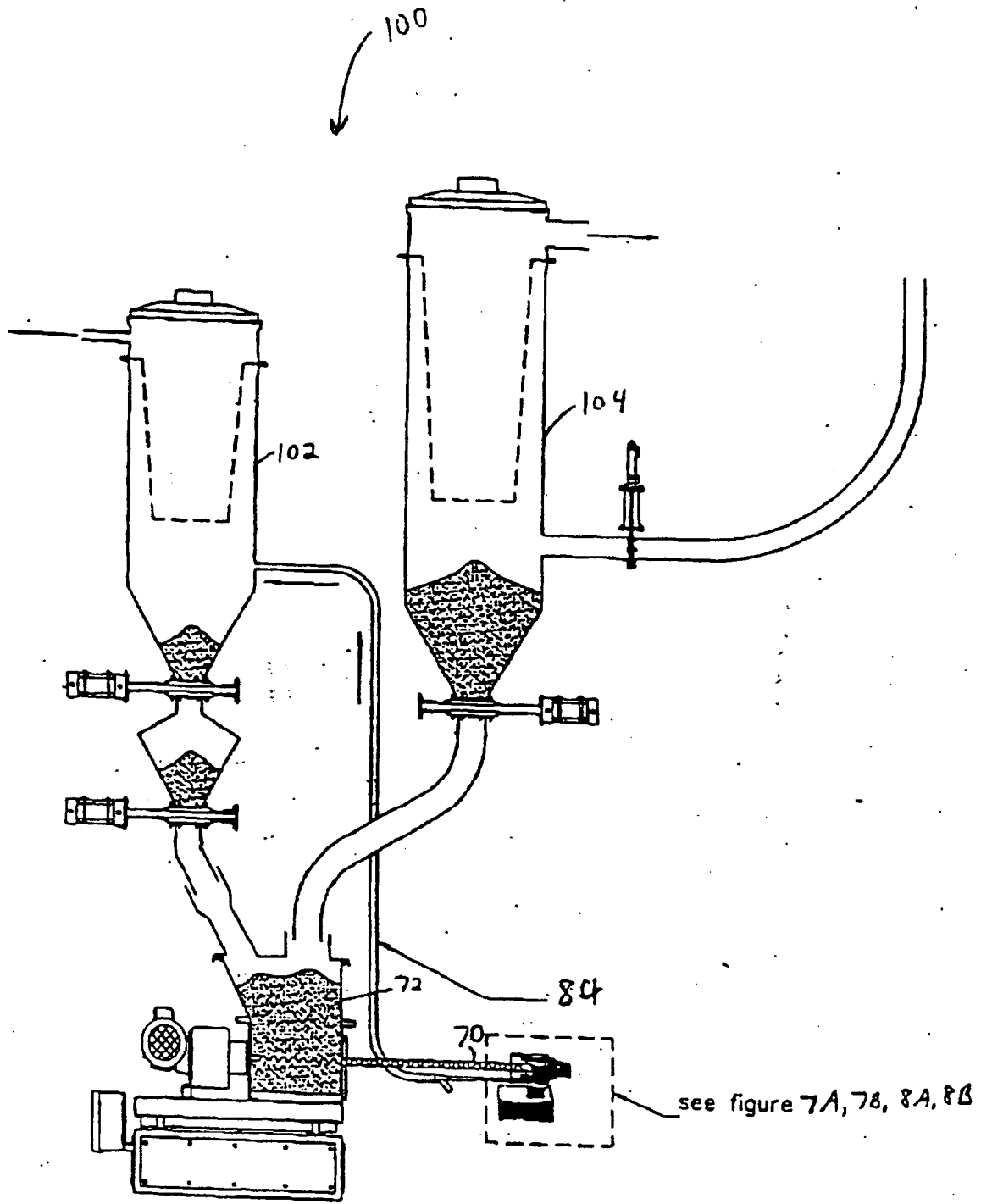


figure 10



GRAVIMETRIC FEEDER

figure 9.



(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 023 884 A3

(12)

## EUROPEAN PATENT APPLICATION

(88) Date of publication A3:

22.11.2000 Bulletin 2000/47

(51) Int. Cl. 7: A61F 13/15

(43) Date of publication A2:

02.08.2000 Bulletin 2000/31

(21) Application number: 99125811.2

(22) Date of filing: 23.12.1999

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU  
MC NL PT SE

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: 23.12.1998 US 220188

10.05.1999 US 309238

(71) Applicant: McNEIL-PPC, INC.

Skillman, New Jersey 08558 (US)

(72) Inventors:

• Rosenfeld, Leonard G.  
East Windsor, NJ 08520 (US)• Foelsch, Lynn  
Hillsborough, NJ 08876 (US)

(74) Representative:

Groening, Hans Wilhelm, Dipl.-Ing.

BOEHMERT &amp; BOEHMERT

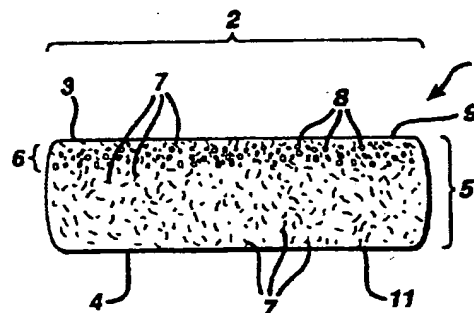
Franz-Joseph-Strasse 38

80801 München (DE)

## (54) Absorbent article with high absorbency zone

(57) The present invention is directed to an absorbent structure comprising an integral absorbent element with an upper surface and a lower surface defining a thickness therebetween. The thickness has an upper 35% containing a high absorbency zone and a lower 65%. The high absorbency zone has a first surface and a second surface separated by a zone thickness. The zone thickness is less than about 35% of the absorbent element thickness. The high absorbency zone comprises absorbent fibers and superabsorbent polymer particles. The superabsorbent polymer particles are present at a loading of no more than 75 grams per square meter and are separated from one another by the absorbent fibers. The lower 65% of the absorbent element thickness is substantially free of superabsorbent polymer particles.

FIG. 1



EP 1 023 884 A3



European Patent  
Office

# EUROPEAN SEARCH REPORT

Application Number  
EP 99 12 5811

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	US 4 186 165 A (ABERSON GERHARD M ET AL) 29 January 1980 (1980-01-29)	1,2,4, 7-9 5,6	A61F13/15
Y	* column 3, line 18,19 * * column 4, line 20 - line 39; claim 2; figures 3,4 *		
X	EP 0 528 248 A (KIMBERLY CLARK CO) 24 February 1993 (1993-02-24) * column 9, line 34 - line 40 * * column 9, line 57 - line 58 * * column 12, line 8 - line 18 * * figure 6 *	1-4,7-9	
X	GB 1 566 594 A (COLGATE PALMOLIVE CO) 8 May 1980 (1980-05-08) * figures 1-3 *	1-4,7-9	
Y	EP 0 719 531 A (KAO CORP) 3 July 1996 (1996-07-03) * abstract; figures 1-6 *	5	TECHNICAL FIELDS SEARCHED (Int.Cl.7)
Y	EP 0 769 284 A (UNI CHARM CORP) 23 Apr11 1997 (1997-04-23) * abstract; figures 1-3 *	6	A61F
The present search report has been drawn up for all claims			
Place of search <b>MUNICH</b>		Date of completion of the search <b>28 September 2000</b>	Examiner <b>Louter, P</b>
<p><b>CATEGORY OF CITED DOCUMENTS</b></p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons &amp; : member of the same patent family, corresponding document</p>			

EPO FORM 1503 03.02 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT  
ON EUROPEAN PATENT APPLICATION NO.**

EP 99 12 5811

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on  
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

28-09-2000

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 4186165	A	29-01-1980	US 4103062 A	25-07-1978
			BE 869988 A	18-12-1978
			ES 245490 U	01-08-1980
EP 0528248	A	24-02-1993	US 5651862 A	29-07-1997
			AU 655047 B	01-12-1994
			AU 2086492 A	18-02-1993
			AU 7441494 A	05-01-1995
			CA 2053964 A	14-02-1993
			DE 69214751 D	28-11-1996
			DE 69214751 T	06-03-1997
			MX 9204120 A	01-02-1993
GB 1566594	A	08-05-1980	ZA 9205014 A	28-04-1993
			AT 366243 B	25-03-1982
			AT 573077 A	15-08-1981
			AU 516668 B	18-06-1981
			AU 2683677 A	11-01-1979
			BE 857608 A	01-12-1977
			BR 7704997 A	02-05-1978
			CA 1092332 A	30-12-1980
			CH 619600 A	15-10-1980
			DE 2734708 A	16-02-1978
			DK 325377 A	10-02-1978
			ES 238917 U	16-02-1979
			FR 2361079 A	10-03-1978
			GR 67256 A	26-06-1981
			IE 45554 B	22-09-1982
			IT 1079991 B	16-05-1985
			JP 1495887 C	16-05-1989
			JP 53021888 A	28-02-1978
			JP 63038442 B	29-07-1988
			MX 148476 A	26-04-1983
			MY 12883 A	31-12-1983
			NL 7708487 A	13-02-1978
			PH 15704 A	14-03-1983
			PT 66812 A, B	01-08-1977
			ZA 7704047 A	28-02-1979
EP 0719531	A	03-07-1996	CN 1134271 A	30-10-1996
			JP 2963647 B	18-10-1999
			JP 8246395 A	24-09-1996
			JP 2955222 B	04-10-1999
			JP 8229070 A	10-09-1996
			US 6086950 A	11-07-2000
			US 5821179 A	13-10-1998

EPO FORM P449

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

**ANNEX TO THE EUROPEAN SEARCH REPORT  
ON EUROPEAN PATENT APPLICATION NO.**

EP 99 12 5811

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on  
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

28-09-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0769284 A	23-04-1997	JP 3053561 B	19-06-2000
		JP 9108262 A	28-04-1997
		AU 702419 B	18-02-1999
		AU 7025496 A	01-05-1997
		BR 9604426 A	23-06-1998
		CA 2188001 A	20-04-1997
		KR 180914 B	01-04-1999
		SG 54375 A	16-11-1998
		US 5795345 A	18-08-1998

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82